Appl. No.

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: June 20, 2003

## AMENDMENTS TO THE CLAIMS

Please cancel Claims 1-7 and 13-15 without prejudice. Please amend Claims 8, 9 and 16, and add new Claims 24-30 as follows:

- 1-7. (Canceled)
- 8. (Currently amended) An isolated polypeptide of <u>comprising</u> arabinose isomerase isolated from *Thermatoga Thermotoga* neapolitana.
- 9. (Currently amended) An isolated polypeptide <u>of-comprising</u> arabinose isomerase encoded by <u>the polynucleotide of Claim 1-a nucleotide derived from Thermotoga neapolitana</u>.
- 10. (Original) The isolated polypeptide of Claim 9, wherein said arabinose isomerase has the amino acid sequence of SEQ. ID NO: 4.
- 11. (Original) The isolated polypeptide of Claim 10, further comprising a solid support.
- 12. (Original) The isolated polypeptide of Claim 11, wherein the solid support is a silica bead.
  - 13-15. (Canceled)
- 16. (Currently amended) An arabinose isomerase produced by the <u>a</u> method of Claim 13. comprising:

providing a host cell transformed with an expression vector comprising a nucleotide derived from *Thermotoga* neapolitana, the polynucleotide coding for an arabinose isomerage; and

culturing the host cell in a medium, thereby producing the arabinose isomerase.

- 17. (Original) A method of producing tagatose, comprising:
  - providing the isolated polypeptide of Claim 9; and
- admixing the arabinose isomerase with galactose, thereby causing a reaction and producing tagatose.
- 18. (Original) The method of Claim 17, wherein the reaction is carried out at a pH from about 5 to about 8.
- 19. (Original) The method of Claim 17, wherein the reaction is carried out at a temperature from about 50°C to about 100°C.
- 20. (Original) The method of Claim 19, wherein the reaction is carried out at a temperature from about 70°C to about 95°C.

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21. (Original) The method of Claim 17, wherein the isolated polypeptide is attached to a solid support.

- 22. (Original) The method of Claim 21, wherein the solid support is a silica bead.
- 23. (Original) The method of Claim 17, wherein the reaction is carried out at a temperature of about 80°C.
- 24. (New) The method of Claim 17, wherein the nucleotide has the sequence of SEQ. ID NO: 3.
- 25. (New) The method of Claim 17, wherein the arabinose isomerase has the amino acid sequence of SEQ. ID NO: 4.
- 26. (New) The isolated polypeptide of Claim 9, wherein the nucleotide has the sequence of SEQ. ID NO: 3.
- 27. (New) The arabinose isomerase of Claim 16, wherein the arabinose isomerase has the amino acid sequence of SEQ. ID NO: 4.
- 28. (New) The arabinose isomerase of Claim 16, wherein the nucleotide has the sequence of SEQ. ID NO: 3.
  - 29. (New) The arabinose isomerase of Claim 16, wherein the host cell is *E. coli*.
- 30. (New) The arabinose isomerase of Claim 16, wherein the host cell is E. coli BL21/DE3 (pTNAI) deposited as Accession No. KCCM-10231.

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**ELECTION OF INVENTION** 

In the Restriction Requirement, the Examiner indicated that this application includes more than one invention identified as follows:

Group I:

Claims 1-7 and 13-15 drawn to DNA, vectors, host cells and

expression of arabinose isomerase;

Group II:

Claim 8-12 and 16 drawn to arabinose isomerase; and

Group III:

Claims 17-23 drawn to a method of producing tagatose.

New Claims 26-30 are drawn to arabinose isomerase and believed to belong to Group II. Applicants elect Group II (Claims 8-12, 16 and 26-30). This election is made without traverse.